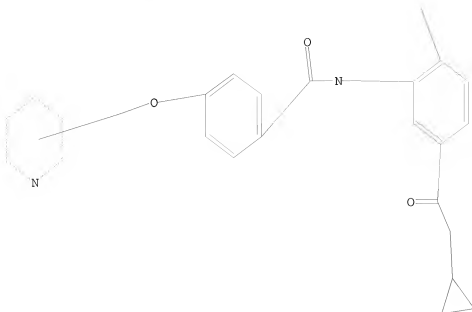


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L1 HAS NO ANSWERS

L1 STR



Structure attributes must be viewed using STN Express query preparation.

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100.0% PROCESSED 7 ITERATIONS

0 ANSWERS

SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**

BATCH **COMPLETE**

PROJECTED ITERATIONS: 7 TO 298

PROJECTED ANSWERS: 0 TO 0

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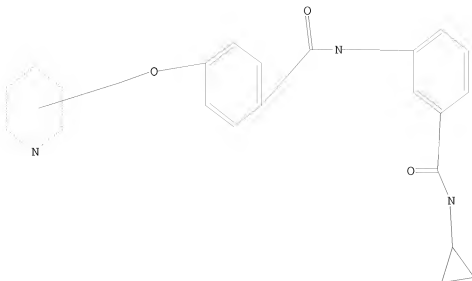
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 L4 HAS NO ANSWERS
 L4 STR



Structure attributes must be viewed using STN Express query preparation.

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100.0% PROCESSED 3005 ITERATIONS 36 ANSWERS
 SEARCH TIME: 00.00.01

L5 36 SEA SSS FUL L4

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 COST IN U.S. DOLLARS SINCE FILE TOTAL
 ENTRY SESSION
 FULL ESTIMATED COST 385.53 385.75

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FILE COVERS 1907 - 24 Feb 2010 VOL 152 ISS 9

FILE LAST UPDATED: 23 Feb 2010 (20100223/ED)
 REVISED CLASS FIELDS (/NCL) LAST RELOADED: Dec 2009
 USPTO MANUAL OF CLASSIFICATIONS THESAURUS ISSUE DATE: Dec 2009

CAPLUS now includes complete International Patent Classification (IPC)
 reclassification data for the third quarter of 2009.

CAS Information Use Policies apply and are available at:

<http://www.cas.org/legal/infopolicy.html>

This file contains CAS Registry Numbers for easy and accurate
 substance identification.

=> s l5

L6 L5

=> d l-2 bib abs hitstr

L6 ANSWER 1 OF 2 CAPLUS COPYRIGHT 2010 ACS on STN
 AN 2009:1260162 CAPLUS

DN 151:433723

TI Pharmaceutical depot comprising N-{5-[(cyclopropylamino)carbonyl]-2-
 methylphenyl}-3-fluoro-4-(pyridin-2-ylmethoxy)benzamide

IN Bateman, Nicola Frances; Macfaul, Philip Alexander; Nash, Ian Alun

PA Astrazeneca AB, Swed.; Astrazeneca UK Limited

SO PCT Int. Appl., 29pp.

CODEN: PIXXD2

DT Patent

LA English

FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
PI	WO 2009125226	A1	20091015	WO 2009-GB50353	20090409
	W:	AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW			
	RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
	US 20090281150	A1	20091112	US 2009-420349	20090408
FRAI	US 2008-43491P	P	20080409		

ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT

AB A pharmaceutical depot is disclosed comprising (i)
 N-{5-[(cyclopropylamino)carbonyl]-2-methylphenyl}-3-fluoro-4-(pyridin-2-ylmethoxy)benzamide, or a pharmaceutically-acceptable salt thereof, as a pharmaceutical agent (PA) and (ii) a polymer which degrades to create an acidic microclimate, wherein the PA is released from the polymer upon polymer degradation

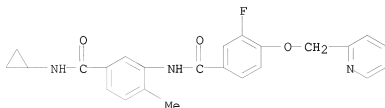
IT 857354-49-5

RL: PEP (Physical, engineering or chemical process); THU (Therapeutic use); BIOL (Biological study); PROC (Process); USES (Uses)

(pharmaceutical depot comprising

N-{5-[(cyclopropylamino)carbonyl]-2-methylphenyl}-3-fluoro-4-(pyridin-2-ylmethoxy)benzamide)

RN 857354-49-5 CAPLUS
 CN Benamide, N-[5-[(cyclopropylamino)carbonyl]-2-methylphenyl]-3-fluoro-4-(2-pyridinylmethoxy)- (CA INDEX NAME)

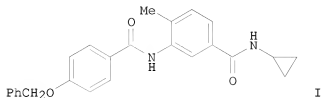


RE.CNT 2 THERE ARE 2 CITED REFERENCES AVAILABLE FOR THIS RECORD
 ALL CITATIONS AVAILABLE IN THE RE FORMAT

L6 ANSWER 2 OF 2 CAPLUS COPYRIGHT 2010 ACS on STN
 AN 2005:588911 CAPLUS
 DN 143:115353
 TI Benamide derivatives bearing a cyclopropylaminoacarbonyl substituent
 useful as cytokine inhibitors
 IN Brown, Dearg Sutherland; Cumming, John Graham; Nash, Ian Alun
 PA AstraZeneca AB, Swed.; AstraZeneca UK Limited
 SO PCT Int. Appl., 88 pp.
 CODEN: PIXXD2
 DT Patent
 LA English
 FAN.CNT 1

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2005061465	A1	20050707	WO 2004-GB5241	20041215
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
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AU 2004303579	B2	20080403		
CA 2547617	A1	20050707	CA 2004-2547617	20041215
EP 1699766	A1	20060913	EP 2004-806056	20041215
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, PL, SK, BA, HR, IS, YU				
CN 1918134	A	20070221	CN 2004-80041887	20041215
BR 2004017844	A	20070417	BR 2004-17844	20041215
JP 2007516979	T	20070628	JP 2006-544544	20041215
NZ 547998	A	20090331	NZ 2004-547998	20041215
MX 2006006660	A	20060811	MX 2006-6660	20060612
ZA 2006004973	A	20070530	ZA 2006-4973	20060615
KR 2006123352	A	20061201	KR 2006-712125	20060619
IN 2006DN03812	A	20070713	IN 2006-DN3812	20060703
NO 2006003330	A	20060914	NO 2006-3330	20060718
US 20070135440	A1	20070614	US 2006-581305	20061012

PRAI GB 2003-29572 A 20031220
 WO 2004-GB5241 W 20041215
 ASSIGNMENT HISTORY FOR US PATENT AVAILABLE IN LSUS DISPLAY FORMAT
 OS CASREACT 143:115353; MARPAT 143:115353
 GI



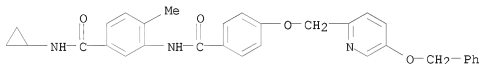
AB The invention concerns a the title compds., or pharmaceutically-acceptable salts; processes for their preparation, pharmaceutical compns. containing them and

their use in the treatment of diseases or medical conditions mediated by cytokines. E.g., I was prepared from 4-benzyloxybenzoic acid and 3-amino-N-cyclopropyl-4-methylbenzamide. Biol. assays include p38 kinase inhibitory, TNF-inhibitory and antiarthritic effects of the compds.

IT 857355-28-3P 857355-36-3P 857355-37-4P
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (benzamide derivs. bearing a cyclopropylaminoacarbonyl substituent useful as cytokine inhibitors)

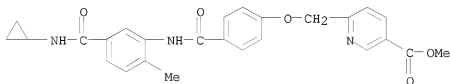
RN 857355-28-3 CAPLUS

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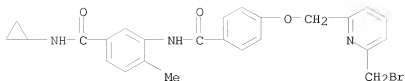
RN 857355-36-3 CAPLUS

CN 3-Pyridinecarboxylic acid, 6-[[4-[[[5-[(cyclopropylamino)carbonyl]-2-methylphenyl]amino]carbonyl]phenoxy]methyl]-, methyl ester (CA INDEX NAME)

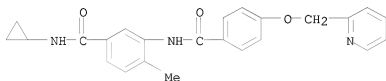


RN 857355-37-4 CAPLUS

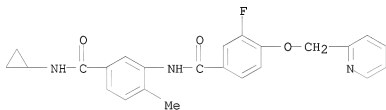
CN Benzamide, 3-[[4-[[6-(bromomethyl)-2-pyridinyl]methoxy]benzoyl]amino]-N-cyclopropyl-4-methyl- (CA INDEX NAME)



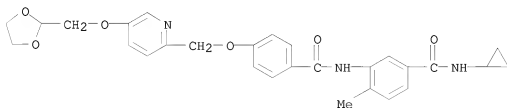
IT 857354-24-6P 857354-49-5P 857354-72-4P
 857355-00-1P
 RL: RCT (Reactant); SPN (Synthetic preparation); THU (Therapeutic use);
 BIOL (Biological study); PREP (Preparation); RACT (Reactant or reagent);
 USES (Uses)
 (benzamide derivs. bearing a cyclopropylaminoacarbonyl substituent
 useful as cytokine inhibitors)
 RN 857354-24-6 CAPLUS
 CN Benzamide, N-cyclopropyl-4-methyl-3-[[4-(2-pyridinylmethoxy)benzoyl]amino]-
 (CA INDEX NAME)



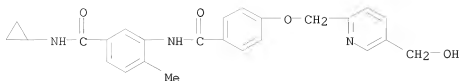
RN 857354-49-5 CAPLUS
 CN Benzamide, N-[5-[(cyclopropylamino)carbonyl]-2-methylphenyl]-3-fluoro-4-(2-
 pyridinylmethoxy)- (CA INDEX NAME)



RN 857354-72-4 CAPLUS
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RN 857355-00-1 CAPLUS
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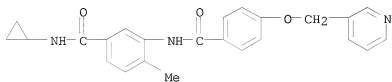
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	857355-08-9P	857355-09-0P	857355-13-6P
	857355-14-7P	857355-15-8P	857355-16-9P
	857355-17-0P		

RL: SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(benzamide derivs. bearing a cyclopropylaminoacarbonyl substituent useful as cytokine inhibitors)

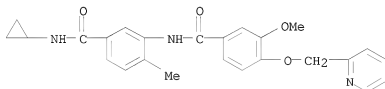
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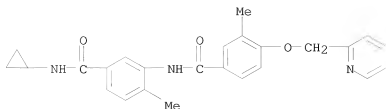
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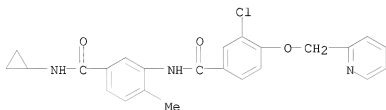
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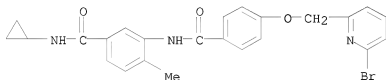
RN 857354-55-3 CAPLUS

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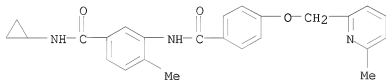
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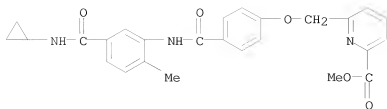
RN 857354-59-7 CAPLUS

CN Benamide, N-cyclopropyl-4-methyl-3-[[4-[(6-methyl-2-pyridinyl)methoxy]benzoyl]amino]- (CA INDEX NAME)



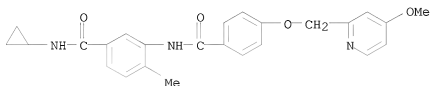
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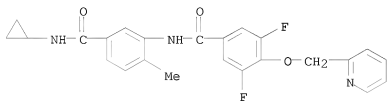
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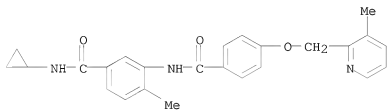
RN 857354-63-3 CAPLUS

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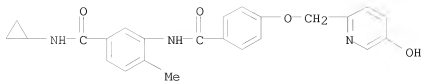
RN 857354-65-5 CAPLUS

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RN 857354-70-2 CAPLUS

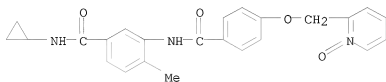
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● HBr

RN 857354-71-3 CAPLUS

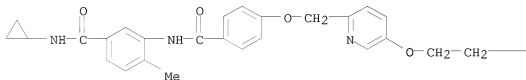
CN Benzamide, N-cyclopropyl-4-methyl-3-[[4-[(1-oxido-2-pyridinyl)methoxy]benzoyl]amino]- (CA INDEX NAME)



RN 857354-73-5 CAPLUS

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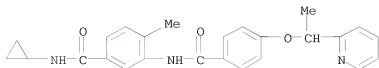


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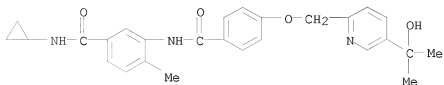
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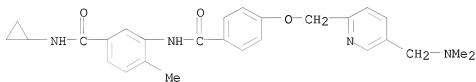
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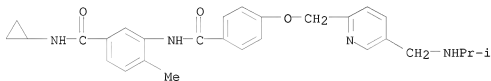
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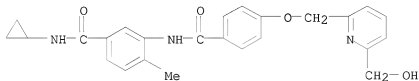
RN 857355-03-4 CAPLUS

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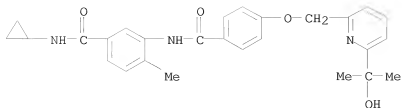
RN 857355-04-5 CAPLUS

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RN 857355-05-6 CAPLUS

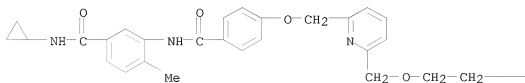
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RN 857355-06-7 CAPLUS

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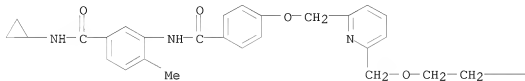
PAGE 1-B

—NEt₂

RN 857355-07-8 CAPLUS

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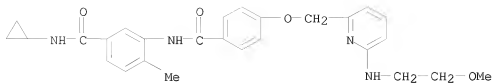


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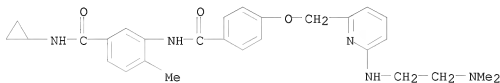
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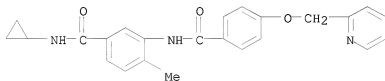
RN 857355-09-0 CAPLUS

CN Benzamide, N-cyclopropyl-3-[[4-[[6-[[2-(dimethylamino)ethyl]amino]-2-pyridinyl]methoxy]benzoyl]amino]-4-methyl- (CA INDEX NAME)



RN 857355-13-6 CAPLUS

CN Benzamide, N-cyclopropyl-4-methyl-3-[[4-(2-pyridinylmethoxy)benzoyl]amino]-, hydrochloride (1:1) (CA INDEX NAME)



● HCl

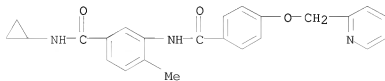
RN 857355-14-7 CAPLUS

CN Benzamide, N-cyclopropyl-4-methyl-3-[[4-(2-pyridinylmethoxy)benzoyl]amino]-, sulfate (1:1) (CA INDEX NAME)

CM 1

CRN 857354-24-6

CMF C24 H23 N3 O3

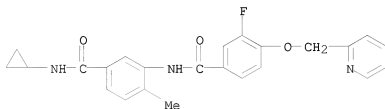


CM 2

CRN 7664-93-9
CMF H2 O4 S

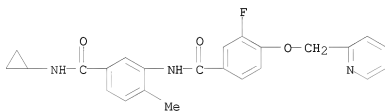


RN 857355-15-8 CAPLUS
CN Benzamide, N-[5-[(cyclopropylamino)carbonyl]-2-methylphenyl]-3-fluoro-4-(2-pyridinylmethoxy)-, hydrochloride (1:1) (CA INDEX NAME)



● HCl

RN 857355-16-9 CAPLUS
CN Benzamide, N-[5-[(cyclopropylamino)carbonyl]-2-methylphenyl]-3-fluoro-4-(2-pyridinylmethoxy)-, hydrobromide (1:1) (CA INDEX NAME)

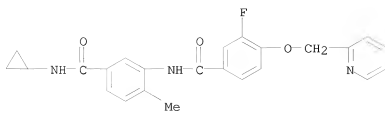


● HBr

RN 857355-17-0 CAPLUS
CN Benzamide, N-[5-[(cyclopropylamino)carbonyl]-2-methylphenyl]-3-fluoro-4-(2-pyridinylmethoxy)-, methanesulfonate (1:1) (CA INDEX NAME)

CM 1

CRN 857354-49-5
CMF C24 H22 F N3 O3



CM 2

CRN 75-75-2

CMF C H4 O3 S

